

## A PHASE 3, MULTICENTER, RANDOMIZED, OPEN-LABEL, STUDY EVALUATING THE EFFICACY AND SAFETY OF NANVURANLAT IN PATIENTS WITH PREVIOUSLY TREATED ADVANCED BILIARY TRACT CANCER

**Status:** Recruiting

### Eligibility Criteria

**Sex:** Male or Female

**Age Group:** 18 years and over

This study is NOT accepting healthy volunteers

**Inclusion Criteria:**

- diagnosed with advanced or metastatic biliary tract cancer (including intrahepatic, extrahepatic, or gallbladder cancer) that cannot be removed with surgery - previously received one platinum-based chemotherapy treatment for advanced disease - able to carry out light daily activities - see link to [clinicaltrials.gov](http://clinicaltrials.gov) for complete Inclusion criteria

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**Exclusion Criteria:**

- received certain cancer treatments, radiation therapy, or major surgery too recently before starting the study - active or untreated brain metastases or uncontrolled serious medical conditions - pregnant or breastfeeding - see link to [clinicaltrials.gov](http://clinicaltrials.gov) for complete Exclusion criteria

### Conditions & Interventions

**Interventions:**

Drug: Nanvuranlat, Other: Physician's Best Choice

**Conditions:**

Cancer

**Keywords:**

Clinics and Surgery Center (CSC), Beacon-BTC, Biliary tract cancer

### More Information

**Description:** This study is for people with advanced biliary tract cancer (cancer of the bile ducts or gallbladder) that cannot be removed with surgery or has spread to other parts of the body. Participants must have previously received one standard chemotherapy treatment for advanced disease. Researchers are studying an investigational medication called nanvuranlat to see if it is safe and effective for treating advanced biliary tract cancer. The study will compare nanvuranlat with standard treatment to learn how well it works, monitor its safety, and better understand how it affects the body and the cancer.

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**Phase:** PHASE3

**IRB Number:** STUDY00026827

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